

# NATIONAL SCIENCE ADVISORY BOARD FOR BIOSECURITY



National Institutes of Health Bldg. 31, C Wing, 6<sup>th</sup> Floor, Conference Room 6 9000 Rockville Pike Bethesda, MD

September 28, 2015

## **MEETING AGENDA**

8:30 – 8:50 am	Welcome and Introductions
8:30 – 8:35 am	Opening Remarks Samuel L. Stanley, M.D. NSABB Chair President, Stony Brook University
8:35 – 8:45 am	Introduction of NSABB Voting and Ex Officio Members Christopher J. Viggiani, Ph.D. Executive Director, NSABB National Institutes of Health
	Review of Conflict of Interest Rules Christopher J. Viggiani, Ph.D.
8:45 – 8:50 am	Approval of NSABB Meeting Minutes Samuel L. Stanley, M.D.
8:50 – 9:50 am	Update from the NSABB Working Group
8:50 – 9:20 am	Update from the Working Group on Evaluating Risks and Benefits of GOF Studies Involving Pathogens with Pandemic Potential Joseph Kanabrocki, Ph.D., C.B.S.P.
	Co-chair, NSABB Working Group Associate Vice President for Research Safety Professor of Microbiology University of Chicago
9:20 – 9:50 am	Associate Vice President for Research Safety Professor of Microbiology
9:20 – 9:50 am 9:50 – 10:00 am	Associate Vice President for Research Safety Professor of Microbiology University of Chicago
	Associate Vice President for Research Safety Professor of Microbiology University of Chicago  NSABB Discussion

### Biosecurity Risk Assessment Kavita Berger, Ph.D.

Scientist, Gryphon Scientific

# **Benefit Assessment**

Corey Meyer, Ph.D.

Senior Analyst, Gryphon Scientific

#### 10:45 – 11:30 am **NSABB Discussion**

11:30 – 11:50 am	Public Comments (Session I)
11:50 – 12:15 pm	Break to Get Lunches for Working Lunch
12:15 – 1:30 pm	The Ethical, Legal, and Policy Issues Associated with Gain-of-Function Studies (Working Lunch)

**Format:** Presentation by author of commissioned ethics analysis followed by comments/remarks by 2 panelists; moderated discussion among panelists and NSABB discussion

#### Questions to be considered during the presentations and discussions:

- What values and decision-making frameworks should NSABB consider in moving beyond the risk/benefit assessment in order to formulate policy recommendations on GOF studies involving pathogens with pandemic potential?
- Is there GOF research that should not be funded and conducted? If so, what are the features of such studies and what considerations should guide the identification of GOF studies that might meet such designation?
- After considering risks and benefits, what policy options or oversight strategies might NSABB consider in generating recommendations to the U.S. Government on the funding and conduct of GOF studies involving pathogens with pandemic potential?

#### Moderator: Susan Wolf, J.D.

Member, NSABB
McKnight Presidential Professor of Law, Medicine & Public Policy
Faegre Baker Daniels Professor of Law
Professor of Medicine
University of Minnesota Law School

# Presenter: Michael Selgelid, Ph.D. Director, Center for Human Bioethics Professor of Bioethics School of Philosophical, Historical and International Studies Monash University

#### Panelists:

Rebecca Dresser, J.D.

# Daniel Noyes Kirby Professor of Law Professor of Ethics in Medicine Washington University in St. Louis

#### Eric Meslin, Ph.D.

Director, Indiana University Center for Bioethics Associate Dean and Professor of Bioethics, Indiana University School of Medicine Managing Director, Center for Law, Ethics and Applied Research in Health Information

1:00 – 1:30 pm	NSABB Discussion
1:30 – 1:50 pm	Public Comments (Session II)
1:50 – 2:05 pm	Break
2:05 – 3:50 pm	NSABB Discussion
	Samuel L. Stanley, M.D.
3:50 – 4:00 pm	Concluding Remarks and Adjourn
	Samuel L. Stanley, M.D.